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DIALINDEX(R)
   (c) 2003 The Dialog Corporation plc
*** DIALINDEX search results display in an abbreviated ***
*** format unless you enter the SET DETAIL ON command. ***
   You have 62 files in your file list.
   (To see banners, use SHOW FILES command)
?s (etch?)(15w)(SiOC or SiCO or oxycarbide or carbon(w)doped(4n)(oxide or dioxide))
Your SELECT statement is:
   s (etch?) (15w) (SiOC or SiCO or oxycarbide or carbon(w) doped(4n) (oxide or
dioxide))
           Items
                  File
                     2: INSPEC 1969-2003/Sep W3
               1
                    8: Ei Compendex(R) 1970-2003/Sep W3
                    34: SciSearch(R) Cited Ref Sci 1990-2003/Sep W3
                    94: JICST-EPlus 1985-2003/Sep W3
                    95: TEME-Technology & Management 1989-2003/Sep W2
                    96: FLUIDEX 1972-2003/Sep
               1
                   144: Pascal 1973-2003/Sep W3
### Status: Break Sent.
?b 2,8,94,144;s (etch?)(15w)(SiOC or SiCO or oxycarbide or carbon(w)doped(4n)(oxide or
dioxide))
       01oct03 14:42:15 User264704 Session D160.4
            $2.45 1.225 DialUnits File411
     $2.45 Estimated cost File411
     $0.46 TELNET
     $2.91 Estimated cost this search
    $26.20 Estimated total session cost 4.723 DialUnits
SYSTEM: OS - DIALOG OneSearch
       2:INSPEC 1969-2003/Sep W3
         (c) 2003 Institution of Electrical Engineers
        2: Alert feature enhanced for multiple files, duplicates
removal, customized scheduling. See HELP ALERT.
       8:Ei Compendex(R) 1970-2003/Sep W3
  File
         (c) 2003 Elsevier Eng. Info. Inc.
  File 94:JICST-EPlus 1985-2003/Sep W3
         (c) 2003 Japan Science and Tech Corp (JST)
  File 144: Pascal 1973-2003/Sep W3
         (c) 2003 INIST/CNRS
      Set Items Description
          _ _ _ _ _
          157008 ETCH?
             291 SIOC
             186 SICO
             755 OXYCARBIDE
          900549 CARBON
          261210 DOPED
          649825 OXIDE
          239742 DIOXIDE
              70
                 CARBON (W) DOPED (4N) (OXIDE OR DIOXIDE)
      Sl
                 (ETCH?) (15W) (SIOC OR SICO OR OXYCARBIDE OR
                  CARBON (W) DOPED (4N) (OXIDE OR DIOXIDE))
?rd
...completed examining records
      S2
               4 RD (unique items)
?t s2/full/all
           (Item 1 from file: 2)
2/9/1
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DIALOG(R) File 2: INSPEC

ン

INSPEC Abstract Number: B2002-10-2550F-034

ashing and cleaning on SiOC-H dielectric: Challenge of Title: characterization and main issues

Author(s): Louveau, O.; Louis, D.; Assous, M.; Blanc, R.; Brun, P.; Lamy, S.; Lajoinie, E.

Author Affiliation: STMicroelectronics, Grenoble, France

Journal: Microelectronic Engineering vol.61-62

Publisher: Elsevier,

7381955

Publication Date: July 2002 Country of Publication: Netherlands

CODEN: MIENEF ISSN: 0167-9317

SICI: 0167-9317(200207)61/62L.867:CACS;1-E

Material Identity Number: F621-2002-003

U.S. Copyright Clearance Center Code: 0167-9317/02/\$22.00

Document Number: S0167-9317(02)00486-0

Document Type: Journal Paper (JP) Language: English

Treatment: Practical (P); Experimental (X)

Abstract: During process development of post etch cleaning for dual damascene copper/ SiOC -H structures, two different kinds of new problems were faced. While a good efficiency is obviously still needed, attention must also be given to dielectric modification and photoresist poisoning which can be induced by the cleaning processes. This study examines different ways of characterization of these phenomena. (6 Refs)

Subfile: B

Descriptors: copper; dielectric thin films; hydrogen; integrated circuit interconnections; photoresists; silicon compounds; sputter etching; surface

Identifiers: SiOC-H dielectric; plasma ashing; cleaning; process development; post etch cleaning; dual damascene copper interconnections; dual damascene Cu/SiOC-H structures; dielectric modification; downstream plasma; photoresist poisoning; Cu-SiOC:H; SiOC:H

Class Codes: B2550F (Metallisation and interconnection technology); B2550E (Surface treatment (semiconductor technology)); B2810 (Dielectric materials and properties); B2550G (Lithography (semiconductor technology)) Chemical Indexing:

Cu-SiOC:H int - SiOC:H int - SiOC int - Cu int - Si int - C int - H int -O int - SiOC:H ss - SiOC ss - Si ss - C ss - H ss - O ss - Cu el - H el - H dop (Elements - 1,3,1,4,5)

SiOC:H sur - SiOC sur - Si sur - C sur - H sur - O sur - SiOC:H ss - SiOC ss - Si ss - C ss - H ss - O ss - H el - H dop (Elements - 3,1,4) Copyright 2002, IEE

(Item 1 from file: 8) 2/9/2

DIALOG(R)File 8:Ei Compendex(R)

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E.I. No: EIP03267514594 06423958

Title: Characterization of low-dielectric-constant SiOC thin films deposited by PECVD for interlayer dielectrics of multilevel interconnection Author: Kim, Hyeong Joon; Shao, Qingyi; Kim, Yoon-Hae

Corporate Source: School of Materials Sci./Eng. Seoul National University, Seoul 151-742, South Korea

Source: Surface and Coatings Technology v 171 n 1-3 Jul 1 2003. p 39-45

Publication Year: 2003

ISSN: 0257-8972 Language: English

Document Type: JA; (Journal Article) Treatment: T; (Theoretical); X; (Experimental)

Journal Announcement: 0307W1

Abstract: Low-dielectric-constant (k) SiOC films were deposited on p-type (100) Si substrates by radio frequency (RF) PECVD with a bistrimethylsilylmethane (BTMSM) precursor and oxygen gas. As more carbon was incorporated into the SiOC film, both the film density and refractive index decreased. This is because the decrease in film density on adding carbon in SiOC films resulted from termination of the Si-O bonding network by replacing oxygen atom with hydrocarbon groups. The hading characteristics of low- R SiOC and SiOF films were investigated by Fourier-transform infrared (FTIR) spectroscopy. The Si-O stretching vibration mode in SiOF films shifted to higher wavenumber with increasing fluorine incorporation, while that in SiOC films shifted to lower wavenumber as the carbon content increased. The origin of the low k of SiOC films was also investigated by determining the electronic, ionic and dipolar contributions. It was found that the low k of SiOC films mainly results from a reduction of the ionic contribution due to a decrease in Si-O bonds in the films. Investigating the compatibility with integration processes showed that modification of the conventional process was required to improve the etch characteristics of the SiOC film. Post-deposition treatment of **sioc** films in hydrogen plasma was effective. The results of this study show that SiOC film is very promising low- k material for interlayer dielectrics of multilevel interconnection. copy 2003 Elsevier Science B.V. All rights reserved. 25 Refs.

Descriptors: *Dielectric films; Silicon compounds; Plasma enhanced chemical vapor deposition; Permittivity; Refractive index; Density (optical); Chemical bonds; Ionic conduction in solids; Carbon inorganic compounds; Oxygen; Fluorine; Molecular vibrations; Fourier transform infrared spectroscopy

Identifiers: Interlayer dielectrics

Classification Codes:

708.1 (Dielectric Materials); 804.2 (Inorganic Compounds); 802.2 (Chemical Reactions); 932.3 (Plasma Physics); 741.1 (Light & Optics); 801.4 (Physical Chemistry); 701.1 (Electricity, Basic Concepts & Phenomena); 931.3 (Atomic & Molecular Physics)

(Electric & Magnetic Materials); 804 (Chemical Products Generally); (Chemical Apparatus & Plants; Unit Operations; Unit Processes); 932 (High Energy Physics; Nuclear Physics; Plasma Physics); 701 (Electricity & Magnetism); 741 (Light, Optics & Optical Devices); 801 (Chemistry); 933 (Solid State Physics); 931 (Applied Physics Generally)

70 (ELECTRICAL ENGINEERING, GENERAL); 80 (CHEMICAL ENGINEERING, GENERAL); 93 (ENGINEERING PHYSICS); 74 (LIGHT & OPTICAL TECHNOLOGY)

2/9/3 (Item 1 from file: 94) DIALOG(R) File 94: JICST-EPlus

(c)2003 Japan Science and Tech Corp(JST). All rts. reserv.

JICST ACCESSION NUMBER: 02A0230058 FILE SEGMENT: JICST-E 05098233 Etching characteristics of SiOC film for low-k dielectrics. YAMASHITA T (1); YAMANAKA M (1); SASAKI T (1); TOMOHISA S (2); SAKAMORI S (2); FUJIWARA N (2); MIYATAKE H (2)

(1) Matsushita Electric Industrial Co., Ltd.; (2) Mitsubishi Electric Corp. Proc Int Symp Dry Process, 2001, VOL.1st, PAGE.153-158, FIG.8, TBL.2, REF.4 JOURNAL NUMBER: Y0378BAP

UNIVERSAL DECIMAL CLASSIFICATION: 621.382.002.2

COUNTRY OF PUBLICATION: Japan LANGUAGE: English

DOCUMENT TYPE: Conference Proceeding

ARTICLE TYPE: Original paper MEDIA TYPE: Printed Publication

ABSTRACT: Etching characteristics of SiOC film for low-k dielectric material (k=2.8-3.0) using fluorocarbon plasmas have been studied employing a dual frequency parallel plate reactive ion etching system. Under the conditions optimized for p-TEOS using a CH2F2CF4/O2Ar gas chemistry, the ${\tt SiOC}$ etch rate and selectivity to photoresist were 164 nm/min and 2.2, slightly lower than those of p-TEOS, and a rough surface was formed at the trench bottom. The etching mechanism for SiOC containing methyl group (-CH3) was examined using optical emission spectroscopy and X-ray photoelectron spectroscopy analyses. As a result of discussion, it was found that C2 and CH radicals produced in the plasma cause the rough surface. Introducing a F-rich CHF3/CF4/O2Ar gas chemistry to compensate this effect in the SiOC etching, the bottom roughness was successfully improved and excellent etched profiles of 0.2.MU.m patterns were realized. (author abst.)

DESCRIPTORS: semiconductor process; RIE; silicon compound; dielectric thin

film; permittivity; con; argon; mixed gas; trench deprocessing); surface roughness; cross section; reaction mechanism

BROADER DESCRIPTORS: production process(control); process; plasma etching; dry etching; etching; surface treatment; treatment; carbon group element compound; thin film; membrane and film; dielectrics; dielectric material; material; ratio; aliphatic chlorine compound; aliphatic halogen compound; organohalogene compound; organochlorine compound; aliphatic fluorine compound; organofluorine compound; rare gas; element; third row element; gas; mixture; object; working and processing; surface quality; flatness(property); property; face; mechanism

CLASSIFICATION CODE(S): NCO3030V

2/9/4 (Item 1 from file: 144)
DIALOG(R)File 144:Pascal
(c) 2003 INIST/CNRS. All rts. reserv.

16151326 PASCAL No.: 03-0306189

Characterization of low-dielectric-constant SiOC thin films deposited by PECVD for interlayer dielectrics of multilevel interconnection

HYEONG JOON KIM; SHAO Qingyi; KIM Yoon-Hae

School of Materials Science and Engineering, Seoul National University, Seoul, 151-742, Korea, Republic of

Journal: Surface & coatings technology, 2003, 171 (1-3) 39-45 ISSN: 0257-8972 CODEN: SCTEEJ Availability: INIST-15987; 354000118190760080

No. of Refs.: 25 ref.

Document Type: P (Serial) ; A (Analytic) Country of Publication: Switzerland

Language: English

Low-dielectric-constant (k) SiOC films were deposited on p-type (100) Si substrates by radio frequency (RF) PECVD with a bis-trimethylsilylmethane (BTMSM) precursor and oxygen gas. As more carbon was incorporated into the SiOC film, both the film density and refractive index decreased. This is because the decrease in film density on adding carbon in SiOC films resulted from termination of the Si-O bonding network by replacing oxygen atoms with hydrocarbon groups. The bonding characteristics of low-k SiOC and SiOF films were investigated by Fourier-transform infrared (FTIR) spectroscopy. The Si-O stretching vibration mode in SiOF films shifted to higher wavenumber with increasing fluorine incorporation, while that in SiOC films shifted to lower wavenumber as the carbon content increased. The origin of the low k of SiOC films was also investigated by determining the electronic, ionic and dipolar contributions. It was found that the low k of SiOC films mainly results from a reduction of the ionic contribution due to a decrease in Si-O bonds in the films. Investigating the compatibility with integration processes showed that modification of the conventional process was required to improve the etch characteristics of the SiOC film. Post-deposition treatment of SiOC films in hydrogen plasma was effective. The results of this study show that SiOC film is very promising low-k material for interlayer dielectrics of multilevel interconnection.

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DIALINDEX(R)
   (c) 2003 The Dialog Corporation plc
*** DIALINDEX search results display in an abbreviated ***
*** format unless you enter the SET DETAIL ON command. ***
   You have 62 files in your file list.
   (To see banners, use SHOW FILES command)
?s (selectivity) (15w) (SiOC or SiCO or oxycarbide or carbon(w) doped(4n) (oxide or dioxide
Your SELECT statement is:
   s (selectivity) (15w) (SiOC or SiCO or oxycarbide or
carbon(w)doped(4n)(oxide or dioxide))
           Items
                   File
           ----
                   _ _ _ _
                     8: Ei Compendex(R) 1970-2003/Sep W3
                    34: SciSearch(R) Cited Ref Sci 1990-2003/Sep W3
                   99: Wilson Appl. Sci & Tech Abs 1983-2003/Aug
                   103: Energy SciTec 1974-2003/Sep B1
                   144: Pascal 1973-2003/Sep W3
                   315: ChemEng & Biotec Abs 1970-2003/Aug
                   347: JAPIO Oct 1976-2003/May(Updated 030902)
                   348: EUROPEAN PATENTS 1978-2003/Sep W03
                   349: PCT FULLTEXT 1979-2002/UB=20030925,UT=20030918
       Examined 50 files
                   354: Ei EnCompassLit(TM) 1965-2003/Sep W4
                   399: CA SEARCH(R) 1967-2003/UD=13914
                   654: US Pat.Full. 1976-2003/Sep 30
   12 files have one or more items; file list includes 62 files.
?b 8,144,354;s (selectivity)(15w)(SiOC or SiCO or oxycarbide or carbon(w)doped(4n)(oxid
e or dioxide))
       01oct03 14:37:09 User264704 Session D160.2
                    2.778 DialUnits File411
            $5.56
     $5.56 Estimated cost File411
     $0.92 TELNET
     $6.48 Estimated cost this search
     $6.51 Estimated total session cost
                                           2.933 DialUnits
SYSTEM:OS - DIALOG OneSearch
         8:Ei Compendex(R) 1970-2003/Sep W3
         (c) 2003 Elsevier Eng. Info. Inc.
  File 144: Pascal 1973-2003/Sep W3
         (c) 2003 INIST/CNRS
  File 354:Ei EnCompassLit(TM) 1965-2003/Sep W4
         (c) 2003 Elsevier Eng. Info. Inc.
*File 354: Ei EnCompassLit/Ei EnCompassPat combined usage is
limited to 2 hrs/yr.
      Set Items Description
          ____
                  ______
          115942 SELECTIVITY
             124 SIOC
              84 SICO
             533
                  OXYCARBIDE
          840092
                 CARBON
          136726 DOPED
          508332
                 OXIDE
          223279
                 DIOXIDE
              40 CARBON (W) DOPED (4N) (OXIDE OR DIOXIDE)
      S1
                  (SELECTIVITY) (15W) (SIOC OR SICO OR OXYCARBIDE OR
                  CARBON (W) DOPED (4N) (OXIDE OR DIOXIDE))
...completed examining records
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S2 3 RD (us le items) ?t s2/full/all

2/9/1 (Item 1 from file: 8)
DIALOG(R)File 8:Ei Compendex(R)
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04003789 E.I. No: EIP94121504788

Title: Isomerization of n-heptane on an oxygen-modified molybdenum carbide catalyst

Author: Blekkan, Edd A.; Pham-Huu, Cuong; Ledoux, Marc J.; Guille, Jean

Corporate Source: Universite Louis Pasteur, Strasbourg, Fr

Source: Industrial & Engineering Chemistry Research v 33 n 7 Jul 1994. p 1657-1664

Publication Year: 1994

CODEN: IECRED ISSN: 0888-5885

Language: English

Document Type: JA; (Journal Article) Treatment: A; (Applications); X; (Experimental)

Journal Announcement: 9502W1

Abstract: The isomerization of n-heptane in the presence of hydrogen has been carried out over a molybdenum carbide catalyst modified by an oxygen treatment. Heptane was isomerized selectively to isoheptanes, a reaction which is difficult over traditional bifunctional catalysts due to extensive cracking. The C//7 products were mainly monomethylhexanes, 2-methylhexane and 3-methylhexane, in close to equilibrium ratios. A typical bifunctional catalyst (Pt supported on an acidic zeolite) gave similar isomerization products, but mostly propane and isobutane as the cracked products. The selectivity over the oxidized carbide was found to be a function of pressure but independent of the conversion; increased the hydrogen pressure led to a decrease in the C//7 selectivity. This was found to be different from the Pt/zeolite catalyst, over which the selectivity was a function of the conversion; a high selectivity was only obtained at low conversions. The active carbide-based catalyst was probably an oxycarbide of molybdenum. The results obtained over the oxidized carbide catalyst are discussed in terms of a bond-shift mechanism via a metallocyclobutane intermediate. (Author abstract) Refs.

Descriptors: *Catalysis; Isomerization; Paraffins; Catalyst selectivity; Catalysts; Carbides; Molybdenum compounds; Pressure effects; Hydrogen; Surface treatment

Identifiers: n-Heptane; Molybdenum carbide; Reaction mechanisms; Intermediates

Classification Codes:

802.2 (Chemical Reactions); 804.1 (Organic Components); 804.2

(Inorganic Components); 802.3 (Chemical Operations)

802 (Chemical Apparatus & Plants); 804 (Chemical Products); 803 (Chemical Agents & Basic Industrial Chemicals)

80 (CHEMICAL ENGINEERING)

2/9/2 (Item 1 from file: 354) DIALOG(R)File 354:Ei EnCompassLit(TM)

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675495 EnCompassLit Document No.: 200114575

Active sites of the isomerization of n-butane over oxygen-modified molybdenum carbide and molybdenum oxides

Author: Liebig S.; Doukkali Kh.; Grunert W.; Gerlach T.

Corporate Source: Lehrstuhl fur Technische Chemie, Ruhr-Univesitat Bochum; Berlin-Adlershof Appl. Chem. Inst.

Source: Studies in Surface Science and Catalysis 130/C 2369-2374 (ISSN 0167--2991) (2000)

Language: English

ISSN: 0167--2991 CODEN: SSCTD

Journal Name: Studies in Surface Science and Catalysis

Document Type: JOURNAL ARTICLE

Publication Date: 0000
Ei EnCompassLit Bulletin Headings: CATALYSTS AND CATALYSIS;
CATALYSTS/ZEOLITES; CHEMICAL PRODUCTS AND PROCESSES; PETROLEUM REFINING
AND PETROCHEMICALS; PETROLEUM-PROCESSING CATALYSTS; PURE HYDROCARBONS
Abstract:

Butane isomerization was studied at 623 K over MoO(sub)3, MoO(sub)2, Mo(sub)9O(sub)2(sub)6, Mo(sub)4O(sub)1(sub)1, etc. Interaction of MoO(sub)3 with n-butane/H(sub)2 at 623 K resulted in gradual development of a highly selective isomerization catalyst. The isomerization site survived contact with air under ambient conditions. XRD provided indications of oxycarbide formation, but no correlation with the isomerization selectivity. XPS showed no indication for the presence of oxycarbide after contact with air. With MoO(sub)3, selectivities > 90% were achieved and held over a prolonged reaction period. With triclinic Mo(sub)4O(sub)1(sub)1, high selectivity was observed without pronounced transient behavior. The active site was an oxide structure of high-valence Mo. Spectra, 5 graphs, and 12 references

Index Terms: ACTIVE SITE; AIR; ANALYTICAL METHOD; BRANCHED CHAIN-P;
 *BUTANE-*A; *C4-*AP; CARBON; CATALYSIS; *CATALYST; DIFFRACTION ANALYSIS
 ; EFFICIENCY; ELEMENT-A; GROUP IVA; GROUP VIA; GROUP VIB;
 *HYDROCARBON-*AP; HYDROGEN-A; *HYDROISOMERIZATION; IDE; *ISOMERIZATION;
 MOLECULAR STRUCTURE; MOLYBDENUM; MOLYBDENUM OXIDE; MOLYBDENUM OXIDE,
 MOO3; OPERATING CONDITION; OXYGEN; PHYSICAL PROPERTY; REACTION
 MECHANISM; REACTION TIME; *SATURATED CHAIN-*AP; SELECTIVITY; *SINGLE
 STRUCTURE TYPE-*AP; *STRAIGHT CHAIN-*A; TEMPERATURE; TEMPERATURE 300 TO
 600 C; TRANSITION METAL; VALENCE; X RAY DIFFRACTION ANALYSIS; YIELD
CAS Registry Numbers: *106-97-8*A; 11098-99-0; 11098-99-0 (BT); 1313-27-5;

1333-74-0A; 7439-98-7; 7440-44-0; 7782-44-7

Sets of Linked Terms: 0006

Linked Terms:

106-97-8A; BUTANE-A; C4-A; HYDROCARBON-A; SATURATED CHAIN-A; SINGLE STRUCTURE TYPE-A; STRAIGHT CHAIN-A

11098-99-0; ACTIVE SITE; CATALYST; GROUP VIA; GROUP VIB; IDE; MOLECULAR STRUCTURE; MOLYBDENUM; MOLYBDENUM OXIDE; OXYGEN; TRANSITION METAL; VALENCE

11098-99-0 (BT); 1313-27-5; ACTIVE SITE; CATALYST; GROUP VIA; GROUP VIB; IDE; MOLECULAR STRUCTURE; MOLYBDENUM; MOLYBDENUM OXIDE; MOLYBDENUM OXIDE, MOO3; OXYGEN; TRANSITION METAL; VALENCE

7439-98-7; 7440-44-0; 7782-44-7; ACTIVE SITE; CARBON; CATALYST; GROUP IVA; GROUP VIA; GROUP VIB; IDE; MOLECULAR STRUCTURE; MOLYBDENUM; OXYGEN; TRANSITION METAL; VALENCE

BRANCHED CHAIN-P; C4-P; HYDROCARBON-P; SATURATED CHAIN-P; SINGLE 1333-74-0A; ELEMENT-A; HYDROGEN-A

2/9/3 (Item 2 from file: 354)

DIALOG(R) File 354: Ei EnCompassLit(TM)

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0556735 EnCompassLit Document No.: 4300332

n-Hexane and n-heptane isomerization at atmospheric and medium pressure on MoO3-carbon-modified supported on SiC and (GAMMA)-Al2O3

Author: Cuong P H; Peschiera E; Gallo P del; Ledoux M J

Corporate Source: Universite Louis Pasteur I; Ecoles des Hautes Etudes des Industries Chimiques de Strasbourg

Source: Applied Catalysis A: General (ISSN 0926-860X) V132 N.1 77-96 (11/9/95)

Language: English ISSN: 0926-860X

Publication Date: 951109

Ei EnCompassLit Bulletin Headings: CATALYSTS/ZEOLITES; CHEMICAL PRODUCTS; CHEMICALS-PROCESSING CATALYSTS; HYDROGENATION; PETROLEUM PROCESSES; PETROLEUM REFINING AND PETROCHEM; PETROLEUM-PROCESSING CATALYSTS; PURE HYDROCARBONS

Abstract:

n-Hexane and n-heptane isomerization at atmospheric and medium

pressure on MoO(sub carbon-modified supported on S .gamma.-Al(sub)20(sub)3. The results obtained for n-hexane and n-heptane isomerization at atmospheric and medium (20 bar) $H(\mathrm{sub})$ 2 pressures over MoO(sub)3 supported on high specific surface area SiC (8.9% Mo) and on .gamma.-Al(sub)2O(sub)3 (19.5% Mo) were compared. The active species was a molybdenum oxycarbide formed by decomposition of the hydrocarbon. This species was extremely active on SiC at 350.degree.C, whereas on .gamma.-Al(sub)20(sub)3, the activity was one order of magnitude lower, and 8.9% Mo on Al(sub)20(sub)3 was inactive. This difference is explained by the fact that on SiC the interaction between MoO(sub)3 and support is low and leads to the formation of a more active phase. Increasing the total pressure improves isomerization activity and selectivity , i.e., formation of the oxycarbide phase is strongly dependent on on the reaction conditions. The results lead to the conclusion that isomerization occurs via a metallacyclobutane intermediate with a high rate of methyl shift. No deactivation with time on stream was observed. Tables, spectra, graphs, and 31 references

Index Terms: *ACTIVITY; ALUMINUM; ALUMINUM OXIDE; C6-A; C7-A; CARBON;
 *CATALYST; *CATALYST ACTIVITY; CATALYST SUPPORT; DEACTIVATION;
 DECOMPOSITION; ELEMENT; GROUP IIIA; GROUP IVA; GROUP VIA; GROUP VIB;
 HEPTANE-A; HEXANE-A; HYDROCARBON-A; HYDROGEN; IDE; *ISOMERIZATION;
 MOLYBDENUM; MOLYBDENUM OXIDE; MOLYBDENUM OXIDE, MOO3; NONE; ON STREAM;
 OPERATING CONDITION; OXYGEN; PARTIAL PRESSURE; *PHYSICAL PROPERTY;
 PRESSURE; *REACTION MECHANISM; SATURATED CHAIN-A; SELECTIVITY; SILICON;
 SILICON CARBIDE; SINGLE STRUCTURE TYPE-A; SPECIFIC SURFACE; STRAIGHT
 CHAIN-A; TEMPERATURE; TEMPERATURE 300 TO 600 C; *USE

CAS Registry Numbers: 110-54-3A; 11098-99-0; 1313-27-5; 1344-28-1; 142-82-5A; 409-21-2

Sets of Linked Terms: 0007

Linked Terms:

110-54-3A; 142-82-5A; C6-A; C7-A; HEPTANE-A; HEXANE-A; HYDROCARBON-A; SATURATED CHAIN-A; SINGLE STRUCTURE TYPE-A; STRAIGHT CHAIN-A

11098-99-0; 1313-27-5; 409-21-2; CARBON; CATALYST; GROUP IVA; GROUP VIA; GROUP VIB; IDE; MOLYBDENUM; MOLYBDENUM OXIDE; MOLYBDENUM OXIDE, MOO3; OXYGEN; SILICON; SILICON CARBIDE; USE

409-21-2; CARBON; CATALYST SUPPORT; GROUP IVA; IDE; SILICON; SILICON CARBIDE; USE

1344-28-1; ALUMINUM; ALUMINUM OXIDE; CATALYST SUPPORT; GROUP IIIA; GROUP VIA; IDE; OXYGEN; USE

11098-99-0; 1313-27-5; 1344-28-1; ALUMINUM; ALUMINUM OXIDE; CATALYST; GROUP IIIA; GROUP VIA; GROUP VIB; IDE; MOLYBDENUM; MOLYBDENUM OXIDE; MOLYBDENUM OXIDE, MOO3; OXYGEN; USE

ELEMENT; HYDROGEN DEACTIVATION; NONE